Natural transformations of tensor algebras and representations of combinatorial groups

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Natural linear and coalgebra transformations of tensor algebras are studied. The representations of certain combinatorial groups are given. These representations are connected to natural transformations of tensor algebras and to the groups of the homotopy classes of maps from the James construction to loop spaces. Applications to homotopy theory appear in a sequel [4].

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1 Introduction

This paper has as its motivation a problem coming from classical homotopy theory, namely, the study of the natural maps from loop suspensions to loop spaces. The method we propose for analyzing properties of certain unstable maps departs from classical unstable homotopy theoretical constructions. We apply the homology functor to natural maps from loop suspensions, and obtain certain functorial coalgebra transformations. This approach justifies the primarily goal of the paper, which is the study of the algebra of natural linear transformations of tensor algebra and related groups of natural coalgebra transformations. These algebras and groups are studied by means of combinatorial group theory. We start with an analogue of a non-commutative exterior algebra defined by Cohen [1] and define several new combinatorial algebras as its generalizations. These algebras are then identified with the corresponding algebras of natural linear transformations of tensor algebras. On the other hand, looking at certain subgroups of the group of units of these combinatorial algebras, we recover the Cohen group K_n and obtain its generalizations (see Section 2 for the definitions). We proceed by establishing group isomorphisms between these combinatorial groups and the corresponding groups of natural coalgebra transformations.

Combinatorial groups, on their own, are closely related to the motivating problem, that is, to the group of natural maps from loop suspensions to loop spaces. This connection

is explained in more detail in the sequel to this paper [4]. There we present solutions to some problems in classical homotopy theory concerning natural maps from loop suspensions to loop spaces obtained by applying the algebraic machinery developed in this paper. Related calculations are done by exploiting the rich structure in combinatorial groups.

Before outlining the main results achieved in the paper, let us explaining the geometrical motivation behind doing the algebra by quoting the main results of the successive paper.

Recall that for any pointed space X, the James–Hopf map $H_k: J(X) \longrightarrow J(X^{(k)})$ is combinatorially defined by

$$H_k(x_1x_2...x_n) = \prod_{1 \le i_1 < i_2 < \dots < i_k \le n} (x_{i_1}x_{i_2}...x_{i_k})$$

with right lexicographical order in the product. The n-th fold Samelson product \widetilde{W}_n on X is given by the composite

$$\widetilde{W}_n: X \wedge \ldots \wedge X \stackrel{E \wedge \ldots \wedge E}{\longrightarrow} \Omega \Sigma X \wedge \ldots \wedge \Omega \Sigma X \stackrel{[[,],\ldots,]}{\longrightarrow} \Omega \Sigma X$$

where $E \colon X \longrightarrow \Omega \Sigma X$ is the canonical inclusion and the second map $[[\,,\,],\ldots,]$ is the n-fold commutator. The n-fold Whitehead product W_n on X is defined as the adjoint of the n-fold Samelson product \widetilde{W}_n .

Theorem A Let X be a pointed space with the null homotopic reduced diagonal $\bar{\Delta}: X \longrightarrow X \wedge X$. Then for n > k,

$$J(X^{(n)}) \xrightarrow{\Omega W_n} J(X) \xrightarrow{H_k} J(X^{(k)})$$

is null homotopic if k does not divide n.

The result concerning the Barratt Conjecture can be formulated as follows.

Theorem B Let $X = \Sigma X'$ be a suspension and let $f: X \longrightarrow \Omega Y$ be a map such that $p^r[f] = 0$ in the group $[X, \Omega Y]$. Let $J(f): J(X) \longrightarrow \Omega Y$ be the canonical multiplicative extension of f. Then the following hold.

- (1) The map $J(f)|_{J_n(X)}: J_n(X) \longrightarrow \Omega Y$ has order p^{r+t} in $[J_n(X), \Omega Y]$ if $n < p^{t+1}$.
- (2) The composite $J_{p^{t+1}}(X) \xrightarrow{J(f)|_{J_{p^{t+1}}(X)}} \Omega Y \xrightarrow{p^{r+t}} \Omega Y$ is homotopic to the composite

$$J_{p^{t+1}}(X) \xrightarrow{\text{pinch}} X^{(p^{t+1})} \xrightarrow{p^{r-1}(\sum_{\tau \in \Sigma_{p^{t+1}-1}} 1 \wedge \tau)} X^{(p^{t+1})} \xrightarrow{\widetilde{W}_{p^{t+1}}} J(X) \xrightarrow{J(f)} \Omega Y,$$

where $p^{r+t}: \Omega Y \longrightarrow \Omega Y$ is the p^{r+t} -th power map, \widetilde{W}_n is the n-fold Samelson product and

$$1 \wedge \tau(x_1 \wedge \cdots \wedge x_{n^{t+1}}) = x_1 \wedge x_{\tau(2)} \wedge \cdots \wedge x_{\tau(n^{t+1})} \colon X^{(p^{t+1})} \longrightarrow X^{(p^{t+1})}$$

is the map which permutes positions.

(3) Let $g = J(f) \circ \widetilde{W}_{p^{t+1}} \circ (\sum_{\tau \in \Sigma_{p^{t+1}-1}} 1 \wedge \tau) \circ p^{r-1} \colon X^{(p^{t+1})} \longrightarrow \Omega Y$. Then g is an equivariant map with respect to the symmetric group action, that is,

$$g \circ \sigma \simeq g$$
 for any $\sigma \in \Sigma_{p^{t+1}}$.

Now we are ready to outline the main results in this paper.

Through the course of the paper R will be a commutative ring with identity unless specified differently. By $\operatorname{Hom}_R(C,A)$ we denote the algebra of natural linear transformations from a coalgebra C to an algebra A, with the multiplication given by the convolution product. The *convolution product* f * g of $f,g: C \longrightarrow A$ is defined by

$$C \xrightarrow{\psi} C \otimes C \xrightarrow{f \otimes g} A \otimes A \xrightarrow{\mu} A$$

where $\psi: C \longrightarrow C \otimes C$ is the comultiplication and $\mu: A \otimes A \longrightarrow A$ is the multiplication.

Let *V* be a free *R*-module. The *James (coalgebra) filtration* $\{J_n(V)\}_{n\geq 0}$ of the tensor algebra T(V) is defined by

$$(1-1) J_n(V) = \bigoplus_{j \le n} T_j(V)$$

for $n \ge 0$, where $T_j(V) = V^{\otimes j}$, the j^{th} stage of the tensor length filtration for T(V). Let $C(V) = J_1(V)$.

A non-commutative analogue of an exterior algebra is given by $A^R(y_1, y_2, \dots, y_n)$ the quotient algebra of the tensor algebra $T(y_1, \dots, y_n)$ over R modulo the two sided ideal generated by the monomials $y_{i_1} \cdots y_{i_t}$ with $i_p = i_q$ for some $1 \le p < q \le t$.

Proposition 1.1 There is an isomorphism of algebras

$$\theta_n: A^R(y_1, \cdots, y_n) \longrightarrow \operatorname{Hom}_R(C(-)^{\otimes n}, T(-)).$$

Furthermore, the James filtration $\{J_n(V)\}_{n\geq 0}$ induces a cofiltration of algebras

$$\operatorname{Hom}_R(T(-), T(-)) \to \cdots \to \operatorname{Hom}_R(J_n(-), T(-)) \to \cdots \to \operatorname{Hom}_R(J_0(-), T(-))$$

where the algebra $\operatorname{Hom}_R(T(-), T(-))$ is given by the inverse limit

$$\operatorname{Hom}_R(T(-), T(-)) \cong \lim_n \operatorname{Hom}_R(J_n(-), T(-)).$$

Let L_n^R be the equalizer of the projection maps

$$\pi_i: A^R(y_1, \cdots, y_n) \longrightarrow A^R(y_1, \cdots, y_{n-1})$$

for $1 \le j \le n$. Then we have the following result.

Proposition 1.2 The map

$$\theta_n = \theta_n \mid_{I^R} : L_n^R \longrightarrow \operatorname{Hom}_R(J_n(-), T(-))$$

is an algebra isomorphism for $n \geq 0$.

For two *R*-modules *C* and *D*, define their *smash product* $C \wedge D$ to be the quotient module

$$C \wedge D = (C \otimes D)/(C \otimes_R R \oplus R \otimes_R D).$$

Proposition 1.3 There are combinatorial algebras $A_n^R[k]$, ${}^RL_n^{(l)}$ and ${}^RL_n^{(l),(k)}$ (for their definitions see Section 3) such that there are algebra isomorphisms

- (1) $\operatorname{Hom}_R(C(-)^{\otimes n}), T(-^{\wedge k})) \cong A_n^R[k]$
- (2) $\operatorname{Hom}_R(J_n(-^{\otimes l}), T(-)) \cong {}^RL_n^{(l)}$
- (3) $\operatorname{Hom}_R(J_n(-^{\otimes l}), T(-^{\otimes k})) \cong {}^RL_n^{(l),(k)}$ for $1 \leq n \leq \infty$.

Let $\operatorname{Coalg}(C, D)$ denote the group of natural coalgebra transformations from a coalgebra C to a Hopf algebra D with the multiplication given by the convolution product. The James filtration $\{J_n(-)\}_{n\geq 0}$ induces a cofiltration of the progroup $\operatorname{Coalg}(T(-), T(-))$. Recall that $C(-) = J_1(-)$.

In [6], Selick and Wu described some properties of the groups $\operatorname{Coalg}(T(-), T(-))$ and $\operatorname{Coalg}(C(-)^{\otimes n}, T(-))$. By the following theorems, we extend their results identifying the groups of natural coalgebra transformations of the James filtration $\{J_n(-)\}_{n\geq 0}$ and their new generalizations with combinatorial groups introduced in Section 1.

Theorem 1.4 There is an isomorphism of groups

$$e: K_n^R \longrightarrow \operatorname{Coalg}(C(-)^{\otimes n}, T(-)) \quad \text{for } n \geq 0.$$

Define \mathcal{H}_n^R to be the equalizer of the projection homomorphisms

$$p_i \colon K_n^R \longrightarrow K_{n-1}^R \quad \text{for } 1 \le j \le n.$$

Theorem 1.5 There is an isomorphism of groups

$$e: \mathcal{H}_n^R \longrightarrow \operatorname{Coalg}(J_n(-), T(-)) \quad \text{for } 1 \leq n \leq \infty.$$

Having in mind the problem solved in Theorem A, we define a generalisation $K_n^R(k)$ of K_n . We set an algebraic notation which is motivated by geometry. Let $\{x_{i_1}|x_{i_2}|\cdots|x_{i_k}\}$ be a notation for a word of length k, in letters $x_{i_1}, x_{i_2}, \ldots, x_{i_k}$. In a successive paper these words will be related to the composite

$$X^n \stackrel{p_{i_1\cdots i_k}}{\longrightarrow} X^{(k)} \stackrel{E}{\longrightarrow} J(X^{(k)}).$$

Let G be a set consisting of all the words $\{x_{i_1}|x_{i_2}|\cdots|x_{i_k}\}$ with $1 \le i_j \le n$ for $1 \le j \le k$ such that $i_s \ne i_t$ for all $1 \le s < t \le k$.

The group $K_n^R(k)$ is defined combinatorially for any commutative ring R so that the generators are elements of G and a certain set of relations, which will be discussed in detail later on in the paper. For k = 1, we denote $K_n^R(1)$ by K_n^R . We will call $K_n^R(k)$ a *Cohen group* as it is a generalization of the combinatorial group $K_n = K_n^{\mathbb{Z}}(1)$ defined by Cohen [1].

Theorem 1.6 There are combinatorial groups $K_n^R(k)$, ${}^R\mathcal{H}_n^{(l)}$ and ${}^R\mathcal{H}_n^{(l),(k)}$ (see Section 1 for their definitions) such that there are group isomorphisms

- (1) $\operatorname{Coalg}(C(-)^{\otimes n}), T(-^{\wedge k})) \cong K_n^R(k)$
- (2) $\operatorname{Coalg}(J_n(-^{\otimes l}), T(-)) \cong {}^R\mathcal{H}_n^{(l)}$
- (3) $\operatorname{Coalg}(J_n(-^{\otimes l}), T(-^{\otimes k})) \cong {}^R\mathcal{H}_n^{(l),(k)}$ for $1 \leq n \leq \infty$.

The compositions of the group isomorphisms in Theorems 1.4, 1.5 and 1.6, with the canonical inclusion of the natural coalgebra transformations Coalg(C, D) of tensor algebras into the natural linear transformations $Hom_R(C, D)$ of tensor algebras give rise to faithful representations of the introduced combinatorial groups to the corresponding algebras of natural linear transformations of tensor algebras.

In the sequel to this paper [4] we establish a connection between the combinatorial groups \mathcal{H}_n^R , ${}^R\mathcal{H}_n^{(I)}$ and ${}^R\mathcal{H}_n^{(I)(k)}$ and the groups of the homotopy classes of maps from the topological James construction J(X) of spaces X with the null homotopic reduced diagonal. We do that by first restricting the ring R to \mathbb{Z} or \mathbb{Z}/p^r and then constructing injective maps:

$$e_X: \mathcal{H}_n^R \longrightarrow [J_n(X), J(X)]$$

$$e_X: {}^R\mathcal{H}_n^{(l)} \longrightarrow [J_n(X^{(l)}), J(X)]$$

$$e_X: {}^R\mathcal{H}_n^{(l)(k)} \longrightarrow [J_n(X^{(l)}), J(X^{(k)})]$$

The disposition of the paper is as follows. Section 2 catalogues all the combinatorial groups of our study and states various properties they satisfy. Section 3 relates combinatorial algebras to natural linear transformations of tensor algebras. Section 4 builds up to and deals with the primary focus of the paper, that is, establishing group isomorphisms between the combinatorial groups defined in Section 2 and the corresponding groups of functorial coalgebra transformations of tensor algebras. Section 5 gives a representation of the combinatorial group $K_n^R(k)$ and relates that group to the group of certain functorial coalgebra transformations.

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2 Cohen groups and their generalizations

In [2, 1], Cohen defined the combinatorial group $K_n^R(x_1, x_2, \dots, x_n)$ for $R = \mathbb{Z}$ or \mathbb{Z}/p^r . Following his approach we define the Cohen group $K_n^R(x_1, x_2, \dots, x_n)$ for any commutative ring R with identity. Let R_x denote a copy of R labeled by x. Write x^r for the element $r \in R_x = R$, and just x for x^1 .

Definition 2.1 The *Cohen group* $K_n^R(x_1, x_2, ..., x_n)$ is the quotient group of the free product $F_n^R = \coprod_{i=1}^n R_{x_i}$ modulo the relations

- (1) $[[x_{i_1}^{r_1}, x_{i_2}^{r_2}], x_{i_3}^{r_3}], \dots, x_{i_l}^{r_l}] = 1$ if $i_s = i_t$ for some $1 \le s, t \le l$;
- (2) $[[x_{i_1}^{r_1}, x_{i_2}^{r_2}], x_{i_3}^{r_3}], \dots, x_{i_l}^{r_l}] = [[x_{i_1}^{r_1'}, x_{i_2}^{r_2'}], x_{i_3}^{r_3'}], \dots, x_{i_l}^{r_l'}]$ if there is an equality $r_1 \cdot r_2 \cdot \dots \cdot r_l = r_1' \cdot r_2' \cdot \dots \cdot r_l'$.

where $[x, y] = x^{-1}y^{-1}xy$ and $[[y_1, y_2], y_3], \dots, y_l]$ is an iterated commutator.

For brevity of notation, we write K_n^R for the Cohen group $K_n^R(x_1, x_2, \ldots, x_n)$ when the generators are assumed, and for $K_n^{\mathbb{Z}}$ we use Cohen's notation K_n . As we will show in the sequel, the Cohen group K_n is closely related to the group $[X^n, J(X)]$ for any space X such that its reduced diagonal $\bar{\Delta}: X \longrightarrow X \wedge X$ is null homotopic.

Define group homomorphisms $p_j: F_n^R \longrightarrow F_{n-1}^R$ and $s_j: F_{n-1}^R \longrightarrow F_n^R$ by

$$p_j(x_i^r) = \begin{cases} x_i^r & \text{for } i < j \\ 1 & \text{for } i = j \\ x_{i-1}^r & \text{for } i > j \end{cases} \qquad s_j(x_i^r) = \begin{cases} x_i^r & \text{for } i < j \\ x_{i+1}^r & \text{for } i \ge j \end{cases}$$

for $1 \leq j \leq n$. It follows that the composites $F_n^R \xrightarrow{p_j} F_{n-1}^R \longrightarrow K_{n-1}^R$ and $F_{n-1}^R \xrightarrow{s_j} F_n^R \longrightarrow K_n^R$ factor through K_n^R and K_{n-1}^R , respectively, inducing the homomorphisms of the Cohen groups $p_j \colon K_n^R \longrightarrow K_{n-1}^R$ and $s_j \colon K_{n-1}^R \longrightarrow K_n^R$.

The next objective is to generalise the Cohen groups K_n , by defining new combinatorial groups related to the group $[X^n, J(X^{(k)})]$.

Definition 2.2 The group $K_n^R(k)$ is defined combinatorially as the quotient group of the free product

$$\prod_{\substack{1 \leq i_j \leq n \\ 1 \leq j \leq k}} R_{\{x_{i_1} | x_{i_2} | \cdots | x_{i_k}\}}$$

modulo the relations given by the following identities:

- (1) $\{x_{i_1}|x_{i_2}|\cdots|x_{i_k}\}^r = 1$ if $i_s = i_t$ for some $1 \le s < t \le k$;
- (2) $[[\{x_{i_1}|x_{i_2}|\cdots|x_{i_k}\}^{r_1},\{x_{i_{k+1}}|x_{i_{k+2}}|\cdots|x_{i_{2k}}\}^{r_2},\cdots \\ \cdots,\{x_{i_{(l-1)k+1}}|x_{i_{(l-1)k+2}}|\cdots|x_{i_{lk}}\}^{r_l}] = 1$ if $i_s = i_t$ for some $1 \le s < t \le kl$, where $[[a_1,a_2,\cdots,a_l] = [[a_1,a_2],a_3],\cdots,a_l]$ with $[x,y] = x^{-1}y^{-1}xy$;

$$(3) \quad [[\{x_{i_{1}}|x_{i_{2}}|\cdots|x_{i_{k}}\}^{r_{1}},\{x_{i_{k+1}}|x_{i_{k+2}}|\cdots|x_{i_{2k}}\}^{r_{2}},\cdots, \\ \qquad \cdots,\{x_{i_{(l-1)k+1}}|x_{i_{(l-1)k+2}}|\cdots|x_{i_{lk}}\}^{r_{l}}] = \\ [[\{x_{i_{1}}|x_{i_{2}}|\cdots|x_{i_{k}}\}^{r'_{1}},\{x_{i_{k+1}}|x_{i_{k+2}}|\cdots|x_{i_{2k}}\}^{r'_{2}},\cdots, \\ \qquad \cdots,\{x_{i_{(l-1)k+1}}|x_{i_{(l-1)k+2}}|\cdots|x_{i_{lk}}\}^{r'_{l}}] \\ \text{if } r_{1} \cdot r_{2} \cdot \ldots \cdot r_{l} = r'_{1} \cdot r'_{2} \cdot \ldots \cdot r'_{l}.$$

Remark In a similar way as for the Cohen group K_n^R , whenever $R = \mathbb{Z}$ we will denote $K_n^R(k)$ by $K_n(k)$. It is obvious that for k = 1, $K_n(1)$ is the Cohen group K_n .

Let q be an integer. Notice that the group $K_n^{\mathbb{Z}/q}(k)$ is the quotient group of $K_n(k)$ modulo the following additional relations:

$$\{x_{i_1}|x_{i_2}|\cdots|x_{i_k}\}^q=1$$

for each generator $\{x_{i_1}|x_{i_2}|\cdots|x_{i_k}\}$.

Selick and Wu [6] defined combinatorial groups \mathcal{H}_n^R in order to study the group $[J_n(X), J(X)]$ with the main goal of obtaining information on the group [J(X), J(X)]. It should be remarked that the concept (and notation) of \mathcal{H}_n^R was invented by Cohen. Here we recall their definition.

Definition 2.3 The group \mathcal{H}_n^R is defined to be the equalizer of the projections $p_j \colon K_n^R(x_1, x_2, \dots, x_n) \longrightarrow K_{n-1}^R(x_1, x_2, \dots, x_{n-1})$ for $1 \le j \le n$.

By definition, as $p_i \mid_{\mathcal{H}_n^R} = p_j \mid_{\mathcal{H}_n^R}$ for $1 \leq i, j \leq n$, there is a homomorphism $d_n \colon \mathcal{H}_n^R \longrightarrow \mathcal{H}_{n-1}^R$ such that the diagram

$$\mathcal{H}_{n}^{R} \xrightarrow{} K_{n}^{R}$$

$$\downarrow^{d_{n}} \qquad \downarrow^{p_{i}}$$

$$\mathcal{H}_{n-1}^{R} \xrightarrow{} K_{n-1}^{R}$$

commutes for each $1 \le i \le n$.

The following lemma was proved by Selick and Wu [6]. As the proof is illustrative and will be used later on in the paper, we include it here.

Lemma 2.4 The homomorphism $d_n: \mathcal{H}_n^R \longrightarrow \mathcal{H}_{n-1}^R$ is an epimorphism for each n.

Proof By induction on k, we show that $d_{k,n}=d_{k+1}\circ\ldots\circ d_n\colon \mathcal{H}_n^R\longrightarrow \mathcal{H}_k^R$ is an epimorphism for $k\leq n$. Clearly, $d_{1,n}$ is an epimorphism. Suppose that $d_{k-1,n}$ is an epimorphism with k>1 and let $\alpha\in\mathcal{H}_k^R$. Since $d_{k-1,n}\colon\mathcal{H}_n^R\longrightarrow\mathcal{H}_{k-1}^R$ is onto, we assume that α lies in the kernel of $d_k\colon\mathcal{H}_k^R\longrightarrow\mathcal{H}_{k-1}^R$. Let

$$\alpha_{k,n} = \prod_{1 \le i_1 < i_2 < \dots < i_{n-k} \le n} s_{i_{n-k}} s_{i_{n-k-1}} \dots s_{i_1} \alpha \in K_n^R$$

with lexicographic order from the right. Then it is routine to check that $\alpha_{k,n} \in \mathcal{H}_n^R$ with $d_{k,n}(\alpha_{k,n}) = \alpha$ and hence the result.

Lemma 2.4 results in a progroup (a tower of group epimorphisms)

$$\mathcal{H}^R \longrightarrow \cdots \longrightarrow \mathcal{H}_n^R \stackrel{d_n}{\longrightarrow} \mathcal{H}_{n-1}^R \longrightarrow \cdots \longrightarrow \mathcal{H}_1^R$$

where \mathcal{H}^R is the group defined by the inverse limit

$$\mathcal{H}^R = \lim_{d_n} \mathcal{H}_n^R$$
.

We recall an important description of the kernel of $d_n: \mathcal{H}_n^R \longrightarrow \mathcal{H}_{n-1}^R$ that will be used later on in the paper.

Let R be a commutative ring and \bar{V} the free R-module with basis $\{x_1, \ldots, x_n\}$. Then $\mathrm{Lie}^R(n)$ denotes the R-submodule of $\bar{V}^{\otimes n}$ generated by the n-fold commutators $[[x_{\sigma(1)}, x_{\sigma(2)}], \ldots, x_{\sigma(n)}]$ for σ a permutation in the symmetric group Σ_n on n letters.

Theorem 2.5 (Cohen [1]) Let $\Lambda(n)$ be the kernel of the group homomorphism $d_n: \mathcal{H}_n^R \longrightarrow \mathcal{H}_{n-1}^R$. Then $\Lambda(n)$ is isomorphic to $\operatorname{Lie}^R(n)$ for $R = \mathbb{Z}$ or \mathbb{Z}/p^r .

In this paper we go a step further by defining new combinatorial groups and algebras in order to study natural transformations of tensor algebras. In a sequel we translate this algebraic setting into geometry which aims at solutions of certain problems of classical homotopy theory. With this in mind we first define two families of groups ${}^{R}\mathcal{H}_{n}^{(l)}$ and ${}^{R}\mathcal{H}_{n}^{(l),(k)}$ that can be seen as generalizations of \mathcal{H}_{n}^{R} and which will shed some light on the study of the groups $[J_{n}(X^{(l)}),J(X)]$ and $[J_{n}(X^{(l)}),J(X^{(k)})]$.

The projection $p_j: K_n^R(x_1, x_2, \dots, x_n) \longrightarrow K_{n-1}^R(x_1, x_2, \dots, x_{n-1})$ is given by

$$p_j(x_i^r) = \begin{cases} x_i^r & \text{for } i < j\\ 1 & \text{for } i = j\\ x_{i-1}^r & \text{for } i > j \end{cases}$$

for $1 \le j \le l$.

Define the projective homomorphism

(2-1)
$$p_{j+\{1,\ldots,l\}}: K_{ln}^R(x_1,x_2,\ldots,x_{ln}) \longrightarrow K_{l(n-1)}^R(x_1,x_2,\ldots,x_{l(n-1)})$$

by

$$p_{j+\{1,...,l\}}(x_i^r) = \begin{cases} x_i^r & \text{for } i < j+1\\ 1 & \text{for } j+1 \le i \le j+l\\ x_{i-1}^r & \text{for } i > j+l \end{cases}$$

for $0 \le j \le n - 1$.

Definition 2.6 Define ${}^{R}\mathcal{H}_{n}^{(l)}$ to be the equalizer of the projections

$$K_{ln}^{R}(x_{1}, x_{2}, \dots, x_{ln}) \cap \left(\bigcap_{j=0}^{n-1} \left(\bigcap_{i=1}^{l} \operatorname{Ker} p_{jl+i}\right)\right)$$

$$p_{j+\{1,\dots,l\}} \downarrow \downarrow \downarrow$$

$$K_{l(n-1)}^{R}(x_{1}, x_{2}, \dots, x_{l(n-1)}) \cap \left(\bigcap_{j=0}^{n-2} \left(\bigcap_{i=1}^{l} \operatorname{Ker} p_{jl+i}\right)\right)$$

for $0 \le j \le n-1$,

As the group ${}^R\mathcal{H}_n^{(l)}$ is given by the equalizer of the projections $p_{j+\{1,\dots,l\}}$ for $0\leq j\leq n-1$, that is, $p_{i+\{1,\dots,l\}}\mid_{R\mathcal{H}_n^{(l)}}=p_{j+\{1,\dots,l\}}\mid_{R\mathcal{H}_n^{(l)}}$ for $0\leq i,j\leq n-1$, there is a homomorphism $d_n\colon {}^R\mathcal{H}_n^{(l)}\longrightarrow {}^R\mathcal{H}_{n-1}^{(l)}$ such that the diagram

$$\begin{array}{ccc}
R \mathcal{H}_{n}^{(l)} & \longrightarrow K_{ln}^{R} \\
\downarrow d_{n} & & \downarrow^{P_{j+\{1,\dots,l\}}} \\
R \mathcal{H}_{n-1}^{(l)} & \longrightarrow K_{l(n-1)}^{R}
\end{array}$$

commutes for each $1 \le j \le n-1$.

Lemma 2.7 The homomorphism $d_n: {}^R\mathcal{H}_n^{(l)} \longrightarrow {}^R\mathcal{H}_{n-1}^{(l)}$ is an epimorphism for each n.

Proof Noticing that the homomorphism

$$K_{ln}^R \bigcap \left(\bigcap_{j=0}^{n-1} \left(\bigcap_{i=1}^{l} \operatorname{Ker} p_{jl+i} \right) \right) \stackrel{p_{j+\{1,\dots,l\}}}{\longrightarrow} K_{l(n-1)}^R \bigcap \left(\bigcap_{j=0}^{n-2} \left(\bigcap_{i=1}^{l} \operatorname{Ker} p_{jl+i} \right) \right)$$

is an epimorphism for each $0 \le j \le n-1$, the proof follows along the lines of the proof of Lemma 2.4.

Lemma 2.7 results in a progroup

$$^{R}\mathcal{H}^{(l)} \longrightarrow \cdots \longrightarrow ^{R}\mathcal{H}_{n}^{(l)} \stackrel{d_{n}}{\longrightarrow} ^{R}\mathcal{H}_{n-1}^{(l)} \longrightarrow \cdots \longrightarrow ^{R}\mathcal{H}_{1}^{(l)}$$

where ${}^{R}\mathcal{H}^{(l)}$ is the group defined by the inverse limit

$${}^{R}\mathcal{H}^{(l)} = \lim_{d_n} {}^{R}\mathcal{H}_n^{(l)}.$$

Definition 2.8 The group ${}^{R}\mathcal{H}_{n}^{(l),(k)}$ is the subgroup of $K_{ln}^{R}(k)$ given as the equalizer of the projections

$$K_{ln}^{R}(k) \cap \left(\bigcap_{j=0}^{n-1} \left(\bigcap_{i=1}^{l} \operatorname{Ker} p_{jl+i}\right)\right)$$

$$p_{j+\{1,...,l\}} \downarrow .. \downarrow$$

$$K_{l(n-1)}^{R}(k) \cap \left(\bigcap_{j=0}^{n-2} \left(\bigcap_{i=1}^{l} \operatorname{Ker} p_{jl+i}\right)\right)$$

for $0 \le j \le n-1$, where the homomorphisms p_j and $p_{j+\{1,\dots,l\}}$ are induced on the generalized Cohen group $K_{ln}^R(k)$ by the projections $p_{j+\{1,\dots,l\}}$ on K_{nl}^R defined by (2–1).

In an analogous fashion as for \mathcal{H}^R and ${}^R\mathcal{H}^{(l)}$, using the fact that ${}^R\mathcal{H}^{(l),(k)}_n$ is given as the equalizer of the projections $p_{j+\{1,\dots,l\}}$, there are group epimorphisms

$$d_n: {}^{R}\mathcal{H}_n^{(l),(k)} \longrightarrow {}^{R}\mathcal{H}_{n-1}^{(l),(k)}$$

such that the diagram

$$\begin{array}{ccc}
^{R}\mathcal{H}_{n}^{(l),(k)} & \longrightarrow & K_{ln}^{R}(k) \\
\downarrow^{d_{n}} & & \downarrow^{p_{j+\{1,\dots,l\}}} \\
^{R}\mathcal{H}_{n-1}^{(l),(k)} & \longrightarrow & K_{l(n-1)}^{R}(k)
\end{array}$$

commutes for each $1 \le j \le n-1$. Thus there is a progroup

$$R\mathcal{H}^{(l),(k)} \longrightarrow \cdots \longrightarrow R\mathcal{H}_{n}^{(l),(k)} \stackrel{d_{n}}{\longrightarrow} R\mathcal{H}_{n-1}^{(l),(k)} \longrightarrow \cdots \longrightarrow R\mathcal{H}_{1}^{(l),(k)}$$

where ${}^{R}\mathcal{H}^{(l),(k)}$ is the group defined by the inverse limit

$${}^{R}\mathcal{H}^{(l),(k)} = \lim_{d_n} {}^{R}\mathcal{H}_n^{(l),(k)}.$$

This introduces all the combinatorial groups we will consider in the paper.

3 Combinatorial algebras and natural linear transformations of tensor algebras

In this section the ground ring is assumed to be a commutative ring R with identity. For a Hopf algebra H over R, denote the comultiplication by $\psi: H \longrightarrow H \otimes H$;

the multiplication by $\mu \colon H \otimes H \longrightarrow H$; the augmentation by $\epsilon \colon H \longrightarrow R$ and the coaugmentation by $\eta \colon R \longrightarrow H$.

Let V be a free R-module. The *James (coalgebra) filtration* $\{J_n(V)\}_{n\geq 0}$ of the tensor algebra T(V) is defined as an R-module by

$$J_n(V) = \bigoplus_{j \le n} T_j(V)$$

for $n \ge 0$, where $T_j(V)$ is the j^{th} stage of the tensor word length filtration of T(V). An coalgebra structure on the filtration is given by requiring that the elements of V are primitive and then multiplicatively extend to all of $J_n(V)$. With this coalgebra structure $J_n(V)$ is a subcoalgebra of the primitively generated Hopf algebra T(V).

Let C be a (graded) coalgebra and let A be a (graded) algebra. The *convolution product* $f * g \text{ of } f, g \colon C \longrightarrow A$ is defined by

$$C \xrightarrow{\psi} C \otimes C \xrightarrow{f \otimes g} A \otimes A \xrightarrow{\mu} A$$

where $\psi: C \longrightarrow C \otimes C$ is the comultiplication and $\mu: A \otimes A \longrightarrow A$ is the multiplication.

Let $\operatorname{Hom}_R(T(-), T(-))$ denote the set of all functorial R-linear maps from T(V) to itself. The convolution product induces a multiplication in $\operatorname{Hom}_R(T(-), T(-))$ under which $\operatorname{Hom}_R(T(-), T(-))$ becomes an algebra over R. Furthermore, the James filtration

$$J_0(V) \subset J_1(V) \subset \cdots \subset J_n(V) \subset \cdots \subset T(V)$$

induces a cofiltration of algebras

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$$\operatorname{Hom}_R(T(-), T(-)) \to \cdots \to \operatorname{Hom}_R(J_n(-), T(-)) \to \cdots \to \operatorname{Hom}_R(J_0(-), T(-)).$$

The purpose of this section is on the one hand to describe connections between combinatorial algebras and natural linear transformations of tensor algebras, and on the other hand to establish the context in which an already developed machinery can be utilized to study connections between the combinatorial groups K_n^R , \mathcal{H}_n^R , \mathcal{H}_n^R and $\mathcal{H}_n^{(l)(k)}$ with certain groups of functorial coalgebra transformations of T(-) and $J_n(-)$. We start by recalling Cohen's definition [1] of a non-commutative analogue of the exterior algebra on $\{y_1, y_2, \dots, y_n\}$.

Definition 3.1 The Cohen algebra $A^R(y_1, y_2, \dots, y_n)$ is defined as the quotient algebra of the tensor algebra $T(y_1, \dots, y_n)$ over R modulo the two sided ideal generated by the monomials $y_{i_1} \cdots y_{i_t}$ with $i_p = i_q$ for some $1 \le p < q \le t$.

Remark As in the case of the Cohen group K_n^R , Cohen [1] gave the definition of $A^R(y_1, \dots, y_n)$ for $R = \mathbb{Z}$ or \mathbb{Z}/p^r , while in our case R can be any commutative ring with identity.

The following result, which Cohen stated for $R = \mathbb{Z}$ or \mathbb{Z}/p^r in [1, Theorem 1.3.2], also holds for an arbitrary commutative ring R.

Proposition 3.2 The algebra $A^R(y_1, \dots, y_n)$ is a graded algebra over R and in degree t, $A(y_1, \dots, y_n)_t$ is a free R-module with basis $y_{i_{\sigma(1)}} \dots y_{i_{\sigma(t)}}$ where $\sigma \in \Sigma_t$ acts on $\{1, 2, \dots, t\}$ and $1 \le i_1 < i_2 < \dots < i_t \le n$.

Proof The proof follows directly from the definition of $A^R(y_1, \dots, y_n)$.

Recall that C(V) is defined as $J_1(V)$.

Definition 3.3 Let V be a free R-module. The algebra $A_n(V)$ is defined to be the subalgebra (under convolution) of $\operatorname{Hom}_R(C(V)^{\otimes n}, T(V))$ generated by the elements y_j given by the composites

$$y_i: C(V)^{\otimes n} \xrightarrow{p_j} C(V) \xrightarrow{q} V \xrightarrow{E} T(V),$$

for $1 \le j \le n$, where $p_i : C(V)^{\otimes n} \longrightarrow C(V)$ is given by

$$p_i(x_1 \otimes x_2 \otimes \cdots \otimes x_n) = \epsilon(x_1) \cdots \epsilon(x_{i-1}) x_i \epsilon(x_{i+1}) \cdots \epsilon(x_n),$$

 $q: C(V) \longrightarrow V$ is the projection and $E: V \longrightarrow T(V)$ is the inclusion.

Lemma 3.4 Let V be a connected graded free R-module. Then, in the algebra $A_n(V)$, the following equation holds for the generators y_i :

$$y_{i_1}y_{i_2}\cdots y_{i_t}=0$$

if $i_i = i_k$ for some $j \neq k$.

Proof The lemma can be easily proved by looking at its geometrical realisation. There exists a space $X = \vee S^{n_{\alpha}}$ with $n_{\alpha} \geq 1$ such that $\widetilde{H}_*(X;R) = V$. Thus $H_*(X;R) \cong C(V)$ as coalgebras and $p_j \colon C(V)^{\otimes n} \longrightarrow C(V)$ is given by

$$(p_i)_*: H_*(X^n; R) \longrightarrow H_*(X; R),$$

where $p_j \colon X^n \longrightarrow X$ is the *j*-th projection. Thus $y_{i_1} \cdots y_{i_t}$ is represented by the composite

$$H_*(X^n:R) \xrightarrow{(p_{i_1\cdots i_t})_*} H_*(X^t:R) \xrightarrow{\gamma} \widetilde{H}_*(X^{(t)}:R) = V^{\otimes t} \longrightarrow T(V).$$

where $p_{i_1 \cdots i_t}(x_1, \cdots, x_n) = (x_{i_1}, \cdots, x_{i_t})$ and $\gamma \colon X^t \longrightarrow X^{(t)}$ is the quotient map. Notice that if $i_j = i_k$ for some $j \neq k$, then there exists a map $f \colon X^n \longrightarrow X^{(t-1)}$ such that

$$\bar{\Delta} \circ f = \gamma \circ p_{i_1 \cdots i_t} \colon X^n \longrightarrow X^{(t)}$$

where $\bar{\Delta}: X^{(t-1)} \longrightarrow X^{(t)}$ is some reduced diagonal map. Notice that

$$(\bar{\Delta})_*: \widetilde{H}_*(X^{(t-1)}; R) \longrightarrow \widetilde{H}_*(X^{(t)}; R)$$

is zero since X is a suspension. The assertion of the lemma follows.

Corollary 3.5 Let *V* be a connected graded free *R*-module. Then the map

$$\theta_n: A^R(y_1, \cdots, y_n) \longrightarrow A_n(V) \subseteq \operatorname{Hom}_R(C(V)^{\otimes n}, T(V))$$

given by $\theta_n(y_i) = y_i$ is a well-defined morphism of algebras.

Now we show that there exists a certain connected graded free R-module V such that $\theta_n: A^R(y_1, \dots, y_n) \longrightarrow A_n(V)$ is a monomorphism.

Lemma 3.6 Let V be a free R-module with $\dim(V) = m$. Suppose that $m \ge n$. Then the homomorphism

$$\theta_n: A^R(y_1, y_2, \cdots, y_n) \longrightarrow A_n(V)$$

is a monomorphism.

Proof Let $X = \bigvee^m S^2$ be the wedge of m copies of the 2-sphere S^2 . Suppose that $m \ge n$. Let $V = H_2(X; R)$. A basis for the R-algebra $A^R(y_1, \dots, y_n)_k$ is given by

$$y_{i_{\sigma(1)}}\cdots y_{i_{\sigma(k)}},$$

where (i_1, \dots, i_k) is taken over $1 \le i_1 < \dots < i_k \le n$ and σ runs over all elements in Σ_k . Notice that $\theta_n(y_{i_{\sigma(1)}} \dots y_{i_{\sigma(k)}})$ is represented by the composite

$$H_*(X^n;R) \xrightarrow{p_{i_{\sigma(1)}\cdots i_{\sigma(k)}}} H_*(X^k;R) \xrightarrow{\gamma} \widetilde{H}_*(X^{(k)};R) \equiv V^{\otimes k} \longrightarrow T(V).$$

Let $\{x_1, \dots, x_m\}$ be a basis for $V = H_2(X; R)$. Let $1 \le j_1 < j_2 < \dots < j_k \le n$ and let $z_1, \dots, z_n \in C(V)$ be such that

- (1) $z_p = 1 \text{ if } p \notin \{j_1, \dots, j_k\};$
- (2) $z_{i_s} = x_{i_s}$.

Then

$$\theta_n(y_{i_{\sigma(1)}}\cdots y_{i_{\sigma(k)}})(z_1\otimes z_2\otimes\cdots\otimes z_n)=\begin{cases}0 & \text{for } (i_1,\cdots,i_k)\neq (j_1,\cdots,j_k)\\x_{j_{\sigma(1)}}\cdots x_{j_{\sigma(k)}} & \text{for } (i_1,\cdots,i_k)=(j_1,\cdots,j_k)\end{cases}$$

The assertion follows.

Lemma 3.7 ((Lemma 2.1 in [6])) Let $\phi_V : V^{\otimes n} \longrightarrow V^{\otimes m}$ be a functorial R-linear map for any free R-module V and let x_1, \dots, x_n be n homogeneous elements in V.

(1) If $\dim_R(V) = n = m$, then the element $\phi_V(x_1 \otimes \cdots \otimes x_n)$ belongs to the R-submodule of $V^{\otimes n}$ spanned by the elements

$$x_{\sigma(1)} \otimes \cdots \otimes x_{\sigma(n)}$$

where σ runs through all elements in Σ_n .

(2) If $n \neq m$, then ϕ_V is the zero map.

Let $\operatorname{Hom}_R(C(-)^{\otimes n}, T(-))$ be the set of all functorial R-linear maps from $C(V)^{\otimes n}$ to T(V) with the convolution product. The same object we will be sometimes denoted by $\operatorname{Hom}_R^{\operatorname{funct}}(C(V)^{\otimes n}, T(V))$.

Proposition 3.8 The homomorphism

$$\theta_n: A^R(y_1, \cdots, y_n) \longrightarrow \operatorname{Hom}_R(C(-)^{\otimes n}, T(-))$$

is an isomorphism of algebras.

Proof By Lemma 3.6, there is a free *R*-module *V* such that

$$\theta_n: A^R(y_1, \cdots, y_n) \longrightarrow A_n(V) \subseteq \operatorname{Hom}_R(C(V)^{\otimes n}, T(V))$$

is a monomorphism. Therefore from the following diagram

$$A^{R}(y_{1}, \cdots, y_{n}) \xrightarrow{\theta_{n}} \operatorname{Hom}_{R}(C(-)^{\otimes n}, T(-))$$

$$\downarrow^{ev}_{V}$$

$$\operatorname{Hom}_{R}(C(V)^{\otimes n}, T(V))$$

we have that the homomorphism

$$\theta_n: A^R(y_1, \cdots, y_n) \longrightarrow \operatorname{Hom}_R(C(-)^{\otimes n}, T(-))$$

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is a monomorphism. To show that θ_n is an epimorphism notice that for any V,

$$C(V)^{\otimes n} = (V \oplus R) \otimes \cdots \otimes (V \oplus R) = \bigoplus_{1 \leq i_1 < \cdots < i_t \leq n} V^{\otimes t}$$

and

$$T(V) = \bigoplus_{m=0}^{\infty} V^{\otimes m}.$$

By Lemma 3.7,

$$\operatorname{Hom}_R(C(-)^{\otimes n}, T(-)) = \bigoplus_{1 \leq i_1 < \dots < i_t \leq n} \operatorname{Hom}_R^{\operatorname{funct}}(V^{\otimes t}, V^{\otimes t}) = \bigoplus_{1 \leq i_1 < \dots < i_t \leq n} R(\Sigma_t).$$

Now the assertion follows from Proposition 3.2.

Definition 3.9 The algebra L_n^R is defined to be the equalizer of the homomorphisms

$$\pi_j: A^R(y_1, \cdots, y_n) \longrightarrow A^R(y_1, \cdots, y_{n-1})$$

for $1 \le j \le n$, where the projection map π_i is given by

$$\pi_j(y_k) = \begin{cases} y_k & \text{for } k < j \\ 0 & \text{for } k = j \\ y_{k-1} & \text{for } k > j. \end{cases}$$

By definition of L_n^R , as $\pi_i \mid_{L_n^R} = \pi_j \mid_{L_n^R}$ for $1 \leq i, j \leq n$, the homomorphisms $\pi_j \colon A^R(y_1, \cdots, y_n) \longrightarrow A^R(y_1, \cdots, y_{n-1})$ induce a homomorphism $d_n \colon L_n^R \longrightarrow L_{n-1}^R$ such that the diagram

$$L_n^{RC} \longrightarrow A^R(y_1, \cdots, y_n)$$

$$\downarrow^{d_n} \qquad \qquad \downarrow^{\pi_j}$$

$$L_{n-1}^{R} \longrightarrow A^R(y_1, \cdots, y_{n-1})$$

commutes for $1 \le j \le n$. The algebra L_{∞}^{R} is defined by the inverse limit

$$L_{\infty}^{R} = \lim_{d_n} L_n^{R}$$
.

The next objective is to find a connection between the combinatorial algebra L_n^R for each $n \ge 0$ and related algebras of natural transformations of the tensor algebra T(-).

Consider the homomorphism

$$\theta_n: A^R(y_1, \cdots, y_n) \longrightarrow A_n(V) \subseteq \operatorname{Hom}_R(C(V)^{\otimes n}, T(V)),$$

of Lemma 3.4, where V is a free R-module. Notice that $J_n(V)$ is the coequalizer of the homomorphisms

$$\iota_j : C(V)^{\otimes (n-1)} \longrightarrow C(V)^{\otimes n}$$

for $1 \le j \le n$, where ι_j is the composite

$$C(V)^{\otimes (n-1)} \xrightarrow{\cong} C^{\otimes (j-1)} \otimes R \otimes C(V)^{\otimes (n-j)} \longrightarrow C^{\otimes n}.$$

Let the *i*-th projection $p_i: C(V)^{\otimes n} \longrightarrow C(V)$ be given by

$$p_i(x_1 \otimes \cdots \otimes x_n) = \epsilon(x_1) \cdots \epsilon(x_{i-1}) x_i \epsilon(x_{i+1}) \cdots \epsilon(x_n).$$

Then

$$p_i \circ \iota_j = \begin{cases} p_i & \text{for } i < j; \\ \eta_{C(V)} \circ \epsilon_{C(V)^{\otimes (n-1)}} & \text{for } i = j; \\ p_{i-1} & \text{for } i > j. \end{cases}$$

Thus there is a commutative diagram

$$A^{R}(y_{1}, \dots, y_{n}) \xrightarrow{\theta_{n}} \operatorname{Hom}_{R}(C(V)^{\otimes n}, T(V))$$

$$\downarrow^{\pi_{j}} \qquad \qquad \downarrow^{\iota_{j}^{*}}$$

$$A^{R}(y_{1}, \dots, y_{n-1}) \xrightarrow{\theta_{n-1}} \operatorname{Hom}_{R}(C(V)^{\otimes (n-1)}, T(V)),$$

where π_j is defined in Definition 3.9. Hence there exists a unique homomorphism of algebras

(3-1)
$$\theta_n: L_n^R \longrightarrow \operatorname{Hom}_R(J_n(-), T(-))$$

such that the diagram

$$L_n^R \xrightarrow{\theta_n} \operatorname{Hom}_R(J_n(-), T(-))$$

$$\downarrow \qquad \qquad \downarrow \qquad \qquad \downarrow$$

$$A^R(y_1, \cdots, y_n) \xrightarrow{\theta_n} \operatorname{Hom}_R(C(-)^{\otimes n}, T(-))$$

$$\downarrow^{\pi_j} \qquad \qquad \downarrow^{\iota_j^*}$$

$$A^R(y_1, \cdots, y_{n-1}) \xrightarrow{\theta_{n-1}} \operatorname{Hom}_R(C(-)^{\otimes (n-1)}, T(-))$$

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commutes for every $1 \le j \le n$. Furthermore, the homomorphism (3–1) preserves the projection homomorphisms, that is, there is a commutative diagram of algebras over R

$$L_n^R \xrightarrow{\theta_n} \operatorname{Hom}_R(J_n(-), T(-))$$

$$\downarrow^{d_n} \qquad \qquad \downarrow$$

$$L_{n-1}^R \xrightarrow{\theta_{n-1}} \operatorname{Hom}_R(J_{n-1}(-), T(-)).$$

In the limit, this gives a homomorphism of co-filtered algebras

$$\theta_{\infty}: L_{\infty}^R \longrightarrow \operatorname{Hom}_R(T(-), T(-)).$$

Theorem 3.10 The algebra L_{∞}^R is isomorphic to $\operatorname{Hom}_R(T(-), T(-))$ as cofiltered algebras, that is, the homomorphism

$$\theta_n: L_n^R \longrightarrow \operatorname{Hom}_R(J_n(-), T(-))$$

is an isomorphism of algebras for each $n \ge 0$.

Proof The proof is given by induction on n. The assertion holds trivially in the cases n = 0, 1. Suppose that the assertion holds for n - 1 with $n \ge 2$. By Lemma 3.6, the map

$$\theta_n: A^R(y_1, \cdots, y_n) \longrightarrow \operatorname{Hom}_R(C(-)^{\otimes n}, T(-))$$

is a monomorphism. Thus the homomorphism

$$\theta_n: L_n^R \longrightarrow \operatorname{Hom}_R(J_n(-), T(-))$$

is a monomorphism. Let Γ_n be the kernel of the homomorphism

$$L_n^R \longrightarrow L_{n-1}^R$$

and let $\tilde{\Gamma}_n$ be the kernel of the homomorphism

$$\operatorname{Hom}_R(J_n(-), T(-)) \longrightarrow \operatorname{Hom}_R(J_{n-1}(-), T(-)).$$

Then

$$\Gamma_n = \bigcap_{1 \leq j \leq n} \ker(\pi_j) \cong \Gamma_n(y_1, \cdots, y_n) \cong R(\Sigma_n),$$

where $\Gamma_n(y_1, \dots, y_n)$ is the R-submodule of $T_n(y_1, \dots, y_n)$ spanned by the monomials $y_{\sigma(1)} \cdots y_{\sigma(n)}$ as σ runs through all elements in Σ_n , and $R(\Sigma_n)$ is a group ring. Let $f \in \tilde{\Gamma}_n$, that is, $f_V \colon J_n(V) \longrightarrow T(V)$ is such that $f_V|_{J_{n-1}(V)} \colon J_{n-1}(V) \longrightarrow T(V)$ is

zero. By assertion (2) of Lemma 3.7, there exists a natural map of R-modules $\phi_V \colon V^{\otimes n} \longrightarrow V^{\otimes n}$ such that the diagram

$$J_n(V) \xrightarrow{f_V} T(V)$$

$$\downarrow \qquad \qquad \uparrow$$

$$V^{\otimes n} \xrightarrow{\phi_V} V^{\otimes n}$$

commutes for each V. By assertion (1) of Lemma 3.7, we have

$$\phi_V(x_1\cdots x_n) = \sum_{\sigma\in\Sigma_n} k_{\sigma} x_{\sigma(1)}\cdots x_{\sigma(n)}$$

for some coefficients $k_{\sigma} \in R$. Let

$$z = \sum_{\sigma \in \Sigma_n} k_{\sigma} y_{\sigma(1)} \cdots y_{\sigma(n)} \in \gamma_n \subseteq L_n^R.$$

Then it is routine to check that

$$\theta_n(z) = f$$
.

Thus the map

$$\theta_n: \ \gamma_n \longrightarrow \tilde{\gamma}_n$$

is an epimorphism. The assertion follows from the 5-Lemma.

By Lemma 3.6, we have

Corollary 3.11 Let *V* be a free *R*-module with $dim_R V \ge n$. Then the homomorphism

$$\theta_n: L_n^R \longrightarrow \operatorname{Hom}_R(J_n(V), T(V))$$

is a monomorphism.

The algebra L_{∞}^{R} is called the universal convolution algebra.

Definition 3.12 The algebra $A_n^R[k]$ is defined as the quotient algebra of the tensor algebra generated by the words $\{y_{i_1}|y_{i_2}|\dots|y_{i_k}\}$ with $1 \le i_j \le n$ for $1 \le j \le k$ over R modulo the two sided ideal generated by the monomials

$$\{y_{i_1}|y_{i_2}|\dots|y_{i_k}\}\{y_{i_{k+1}}|y_{i_{k+2}}|\dots|y_{i_{2k}}\}\dots\{y_{i_{(l-1)k+1}}|y_{i_{(l-1)k+2}}|\dots|y_{i_k}\}$$

with $i_s = i_t$ for some $1 \le s < t \le k$.

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a) reaction of alcohols with appropriately substituted 2- or 4- halo- or sulfonate esters of nitrobenzenes or benzonitriles (Method 4A) either neat or in an appropriate solvent such as tetrahydrofuran, dioxane, acetonitrile, N,N-dimethylformamide or dimethylsulfoxide in the presence or absence of one or more molar equivalents of a base such as sodium carbonate, potassium carbonate, sodium hydroxide, potassium hydroxide, sodium hydride, potassium hydride, or the like, at temperatures ranging from room temperature to the reflux temperature of the solvent;

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- b) reactions of preformed sodium, lithium, or potassium phenoxides with
 appropriately substituted 2- or 4- halo- or sulfonate esters of nitrobenzenes or benzonitriles (Method 4H) either neat or in an appropriate solvent such as tetrahydrofuran, dioxane, acetonitrile, N,N-dimethylformamide or dimethylsulfoxide, at temperatures ranging from room temperature to the reflux temperature of the solvent, or;
- c) reaction of ammonia, primary or secondary amines with appropriately substituted 2- or 4-halo- or sulfonate esters of nitrobenzenes or benzonitriles (Methods 4C,F) either neat or in an appropriate solvent such as tetrahydrofuran, dioxane, acetonitrile, N,N-dimethyl-formamide or dimethylsulfoxide, at temperatures ranging from room temperature to the reflux temperature of the solvent;
- d) reaction of preformed sodium, lithium, or potassium salts of amines with appropriately substituted 2- or 4- halo- or sulfonate esters of nitrobenzenes or benzonitriles (Method 4G) in an appropriate solvent such as tetrahydrofuran at temperatures ranging from 0°C to the reflux temperature of the solvent, or;
- e) reaction of sodium sulfide with appropriately substituted 2- or 4- halo- or sulfonate esters of nitrobenzenes or benzonitriles either neat or in an appropriate solvent such as tetrahydro-furan, dioxane, acetonitrile, N,N-dimethylformamide or dimethylsulfoxide, at temperatures ranging from room temperature to the reflux temperature of the solvent, followed by the addition of an alkyl halide directly to the reaction mixture (Method 4E).
- Alternatively, referring to Methods 5C and 6, the nitrobenzene intermediates which are ultimately converted into amines 2, wherein at least one substitutent R₁-R₃ is defined as alkoxy may be prepared from the corresponding substituted hydroxynitrobenzenes by methods which include the following:

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- a) reaction of the hydroxy-nitrobenzene with an alkyl halide or dialkyl sulfonate ester (Method 5C) in the presence of a base, such as potassium carbonate, sodium carbonate, potassium hydroxide, sodium hydroxide, potassium hydride, or sodium hydride, in an appropriate solvent such as acetone, N,N-dimethylformamide, tetrahydrofuran or dimethylsulfoxide at temperatures ranging from room temperature to the reflux temperature of the solvent, or;
- b) reaction of the hydroxy-nitrobenzene with an alkyl alcohol, triphenylphosphine, and a dialkylazadicarboxylate reagent (Method 6), such as diethylazodicarboxylate, in an anhydrous aprotic solvent such as diethyl ether or tetrahydrofuran at temperatures ranging from 0°C to the reflux temperature of the solvent, essentially according to methods described in Mitsunobu, O, Synthesis 1981, 1 and references therein.

In addition, referring to Method 5A and 5E, the carbamoyl amine derivatives utilized as starting materials in Methods 3A-3C which are ultimately converted into amines 2, wherein at least one substituent R₁-R₅ is defined as alkoxy may be prepared the corresponding substituted hydroxy arylamino-tert-butyl-carbamate by reaction with alkyl halides, trifluormethane-sulfonates, 4-methylbenzenesulfonates, dialkylsulfonate, ethylene carbonate and the like in the presence of a suitable base such as potassium carbonate in an appropriate solvent such as acetone, toluene, or N,N-dimethyl-formamide at temperatures ranging from room temperature to the reflux temperature of the solvent.

Alternatively, referring to Methods 7A-G, the nitrobenzene intermediates which are ultimately converted into amines 2, R₁ and/or R₃ is alkoxy, and R₂ and/or R₄ is a halogen, and X equals a bond, may be prepared by standard halogenation reactions which include the following:

- a) reaction of a 2- or 4- hydroxy-nitrobenzene with aqueous sodium hypochlorite (Methods 7A and 7B), at room temperature or;
- b) reaction of a 2-hydroxy-4-methoxy or 2,4-dimethoxynitrobenzene (Method 7C and 7D) with bromine in suitable solvent such as chloroform, dichlormethane, glacial acetic acid or the like in the presence or the absence of silver trifluoroacetate at room temperature, or;

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- c) reaction of a 2,4-dimethoxynitrobenzene (Method 7E) with benzyltrimethylammonium dichloroiodate in the presence of anhydrous zinc chloride in a suitable solvent such as glacial acetic acid, at room temperature or;
- d) reaction of a 2-hydroxy-4-methoxynitrobenzene (Method 7F) with benzyltrimethyl-ammonium dichloroiodate in the presence of sodium bicarbonate in a suitable solvent mixture such as dichloromethane and methanol, at room temperature or;
- e) reaction of a 2,4-dimethoxynitrobenzene (Method 7G) with 3,5-dichloro-1-fluoropyridine triflate in a suitable solvent such as tetrachloroethane, at a temperature ranging from room temperature to the reflux temperature of the solvent.

Referring to Method 8, the nitrobenzene intermediates which are ultimately converted into amines 2, wherein $R_4 = -CF_3$, and R_1-R_3 and R_5-R_8 are defined as above and X equals a bond may be prepared from the corresponding substituted 4-iodonitrobenzenes by reaction with trimethyl(trifluoromethyl)silane in the presence of cuprous iodide and potassium fluoride in a suitable solvent such as N,N-dimethyl-formamide or the like at a temperature ranging from room temperature to the reflux temperature of the solvent in a sealed reaction vessel.

Referring to Methods 19A and 19B, the nitrobenzene intermediates which are ultimately converted into amines 2, wherein $R_4 = -HNCOCH_2NR_7R_8$ or - $HNCOCH_2SR_6$, and R_1-R_3 and R_5-R_8 are defined as above and X equals a bond may be prepared from the corresponding substituted 4-(N-chloroacetyl)-nitroaniline by reaction with either a suitable secondary amine such as dimethylamine, morpholine or the like in a suitable solvent such as tetrahydrofuran and/or water mixtures at temperatures ranging from room temperature to the reflux temperature of the solvent or by reaction with an appropriate thiol in the presence of a suitable base such as sodium or potassium carbonate or the like in a suitable solvent such as tetrahydrofuran, 1,4-dioxane or the like at temperatures ranging from room temperature to the reflux temperature of the solvent.

Referring to Method 25, the nitrobenzene intermediates which are ultimately converted into amines 2, wherein at least one substituent of R_1 - R_5 is defined as triflate and X equals a bond may be prepared from the corresponding phenol by reaction with trifluoromethane sulfonic anhydride in the presence of a tertiary amines such as

triethylamine or diisopropyl-ethylamine or the like in a suitable solvent such as dichloromethane at temperatures ranging from 0°C to room temperature.

Referring to Methods 9, 9B, and 10, the carbamoyl amine derivatives utilized as starting materials in Methods 3A-3C which are ultimately converted into amines 2, wherein at least one substituent R_1-R_5 is defined as either alkylsulfenyl or alkylsulfinyl, may be prepared by reaction of the appropriate 4-alkylthio acylarylamino or carbamoyl arylamino derivative with an appropriate oxidizing agent such as dimethyloxirane or sodium periodate in a suitable solvent mixture such as acetone and dichloromethane or water at room temperature.

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Referring to Method 12, the carbamoyl amine derivatives utilized as starting materials in Methods 3A-3C which are ultimately converted into amines 2, wherein R_4 is defined as 1-hydroxyethyl and R_1 - R_3 and R_5 are defined as above and X equals a bond may be prepared by reacting the corresponding 4-vinyl carbamoyl aniline with sodium borohydride in the presence of mercuric acetate in a suitable solvent such as tetrahydrofuran, 1,4-dioxane or the like and water at room temperature.

Referring to Method 13, the carbamoyl amine derivatives utilized as starting materials in Methods 3A-3C which are ultimately converted into amines 2, wherein R_4 is defined as 2-hydroxyethyl and R_1 - R_3 and R_5 are defined as above and X equals a bond, may be prepared by reacting the corresponding 4-vinyl carbamoyl aniline with sodium borohydride in the presence of glacial acetic acid in a suitable solvent such as tetrahydrofuran, 1,4-dioxane or the like at temperatures ranging from 0°C to room temperature.

Referring to Method 14, the carbamoyl amine derivatives utilized as starting materials in Methods 3A-3C which are ultimately converted into amines 2, wherein R_4 is defined as 1-azidoethyl and R_1 - R_3 and R_5 are defined as above and X is defined above may be prepared by reacting the corresponding 4-(1-hydroxyethyl) carbamoyl aniline with hydrazoic acid in the presence of a dialkylazodicarboxylate such as diethylazodicarboxylate and triphenylphosphine in a suitable solvent mixture such as tetrahydrofuran and dichloromethane at temperatures ranging from 0°C to room temperature.

Referring to Method 15, the carbamoyl amine derivatives utilized as starting materials in Methods 3A-3C which are ultimately converted into amines 2, wherein R_4 is defined as 3-dimethylaminoprop-1-ynyl and R_1 - R_3 and R_5 are defined as above

and X is defined above, may be prepared by reacting the corresponding 4-iodocarbamoyl aniline with 1-dimethylamino-2-propyne in a suitable tertiary amine solvent such as triethylamine or diisopropylethylamine in the presence of bis(triphenylphosphine)palladium(II) chloride and cuprous iodide at temperatures ranging from room temperature to the reflux temperature of the solvent.

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Referring to Method 16, the carbamoyl amine derivatives utilized as starting materials in Methods 3A-3C which are ultimately converted into amines 2, wherein R_4 is defined as 3-dimethylaminoacryloyl and R_1 - R_3 and R_5 are defined as above and X equals a bond, may be prepared by reacting the corresponding 4-(3-dimethylaminoprop-1-ynyl)carbamoyl aniline with a suitable peracid such as 3-chloroperoxybenzoic acid in a suitable solvent mixture such as dichloromethane and methanol at temperatures ranging from 0°C to room temperature.

Referring to Methods 17 and 18, the carbamoyl amine derivatives utilized as starting materials in Methods 3A-3C which are ultimately converted into amines 2, wherein R_4 is defined as either 4-isoxazol-5-yl or 4-(1H-pyrazol-3-yl) and R_1 - R_3 and R_5 are defined as above and X equals a bond, may be prepared by reacting the corresponding 4-(3-dimethylamino-acryloyl)carbamoyl aniline with either hydroxylamine hydrochloride or hydrazine hydrate in a suitable solvent such as 1,4-dioxane or ethanol and the like at room temperature.

Referring to Method 20, the carbamoyl amine derivatives utilized as starting materials in Methods 3A-3C which are ultimately converted into amines 2, wherein $R_4 = -HNCO_2Z$, and R_1-R_3 , R_5 , and Z are defined as above and X equals a bond, may be prepared by reacting the corresponding 4-aminocarbamoyl aniline with 1,1-carbonyl-di-(1,2,4)-triazole and an appropriately substituted alcohol in a suitable solvent mixture such as tetrahydrofuran and dichloromethane and the like at temperatures ranging from room temperature to the reflux temperature of the solvent.

Referring to Methods 26 and 30, the carbamoyl amine derivatives utilized as starting materials in Methods 3A-3C which are ultimately converted into amines 2, wherein at least one substituent of R_1 - R_s is defined as dialkylamino and X is defined above may be prepared by reaction of appropriately substituted aldehydes in the presence of either sodium cyanoboro-hydride or hydrogen gas and 10 % palladium on carbon in a suitable solvent such as water, methanol, tetrahydrofuran mixtures or toluene or the like at room temperature.

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Referring to Methods 27 and 28, amines 2 wherein at least one substituent of R_1 - R_5 is defined as hydroxy and X is defined above can be prepared by reaction of the corresponding ester such as acetate with an appropriate base such as sodium bicarbonate or sodium hydroxide in a suitable solvent mixture such as methanol-water mixtures at temperatures ranging from room temperature to the reflux temperature of the solvent.

Referring to Method 29, amines 2 wherein at least one substituent of R_1 - R_5 is defined as 2-hydroxybenzamido and X is defined above can be prepared by reaction of the corresponding N-(4-aminophenyl)phthalimide with lithium borohydride in an appropriate solvent such as tetrahydrofuran, diethyl ether, or the like at room temperature.

The intermediate amines 2 wherein R_1 - R_5 are defined as above and X equals either - CH_2 - or - $(CH_2)_2$ - can be prepared by the following procedures:

- a) reduction of an appropriately substituted benzo- or phenylacetonitrile with borane-dimethylsulfide complex in a suitable solvent such as ethylene glycol dimethyl ether, tetrahydrofuran or the like a temperatures ranging from room temperature to the reflux temperature of the solvent. (Method 44);
- 11. reduction under one or more atmospheres of hydrogen in the presence of a suitable catalyst such as 5 % or 10 % palladium on carbon and an acid such as 4-methyl-benzenesulfonic acid, hydrochloric acid or the like in a suitable solvent such as ethylene glycol monomethyl ether, ethyl acetate, ethanol or the like at room temperature. (Method 50);
- 12. reduction with lithium aluminum hydride in a suitable solvent such as tetrahydrofuran or diethyl ether at temperatures ranging from 0°C to room temperature. (Method 51);

The unsaturated nitro precursors which are utilized as starting materials in Method 51 and are ultimately converted to amines 2 wherein R_1 - R_5 are defined as above and X equals - $(CH_2)_2$ - can be prepared by reaction of an appropriately substituted benzaldehyde with nitro-methane in the presence of ammonium acetate in a suitable solvent such as acetic acid at temperatures ranging from room temperature to the reflux temperature of the solvent. (Method 53); The benzaldehydes, utilized as starting materials in Method 53, can be prepared by diisobutylaluminum hydride

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reduction of an appropriately substituted benzonitrile. (Method 52) The substituted benzonitriles, utilized as starting materials in Method 52, can be prepared from the corresponding aryl bromide by reaction with copper cyanide in a suitable solvent such as N,N-dimethylformamide at temperatures ranging from room temperature to the reflux temperature of the solvent. (Method 59)

For amines 2, wherein R_1 - R_5 is defined as above and X equals either - $O(CH_2)_2NH_2$ or $-S(CH_2)_2NH_2$, the requisite nitrile precursors may be prepared by reaction of an appropriately substituted phenol or thiophenol with bromoacetonitrile in the presence of a suitable base such as potassium carbonate in an appropriate solvent such as acetone at room temperature according to Method 49.

Alternatively, for amines 2, wherein R_1 - R_5 are defined as above and X equals -(CH_2)₃-, the nitrile precursors can be prepared essentially according to the procedure of Wilk, B. Synthetic Comm. 23, 2481 (1993), by reaction of an appropriately substituted phenethanol with acetone cyanohydrin and triphenylphosphine in the presence of a suitable azodicarboxylate such as diethyl azodicarboxylate in an appropriate solvent such as diethyl ether or tetrahydro-furan or the like at temperatures ranging from 0°C to room temperature. (Method 54)

Alternatively, intermediate amines 2 wherein R₁-R₅ are defined as above and X equals -(CH(CH₃))- can be prepared by acid or base catalyzed hydrolysis of the corresponding formamide using an appropriate acid catalyst such as 6N hydrochloric acid or a suitable base catalyst such as 5N sodium or potassium hydroxide in an appropriate solvent mixture such as water and methanol or water and ethanol at temperatures ranging from room temperature to the reflux temperature of the solvent. (Method 46)

The formamide precursors utilized as starting materials in Method 46 and which are ultimately converted into amines 2, are prepared according to Method 45 by treatment of an appropriately substituted acetophenone with ammonium formate, formic acid and formamide at temperatures ranging from room temperature to the reflux temperature of the solvent.

Alternatively, amines 2 wherein R_1 - R_5 are defined as above and X equals - (CH(CH₃))- can be prepared by reduction of an appropriately substituted O-methyl oxime in the presence of sodium borohydride and zirconium tetrachloride in a suitable solvent such as tetrahydrofuran or diethyl ether at room temperature Method

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48 essentially according to the procedure of Itsuno, S., Sakurai, Y., Ito, K. Synthesis 1988, 995. The requisite O-methyl oximes can be prepared from the corresponding acetophenone by reaction with methoxylamine hydrochloride and pyridine in a suitable solvent such as ethanol or methanol at temperatures ranging from room temperature to the reflux temperature of the solvent. (Method 47)

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Amines 2 for which R₁-R₃ are defined as above and X equals -CH(J)- where J is defined as above, can be prepared by reduction of the appropriately substituted ketone by the methods described above (Methods 45, 47, and 48). These requisite ketones, when not commercially available, can be prepared by reaction of a suitably substituted benzaldehyde with an appropriate organometallic reagent such as phenyllithium, isopropylmagnesium bromide or ethylmagnesium bromide or the like in a suitable solvent such as diethyl ether or tetrahydrofuran at temperatures ranging from -78 °C to 0°C. (Method 57) The resulting alcohols can be oxidized to the corresponding ketone with an appropriate oxidizing agent such as chromium trioxide in aqueous sulfuric acid and acetone or pyridinium chlorochromate or pyridium dichromate in an appropriate solvent such as dichloromethane or the like at room temperature. (Method 58)

The intermediate anilines 5 may be prepared as previously described Method 3A. Thus treating phenyl carbamic acid tert-butyl ester 6, wherein X equals a bond and G are described as above, with neat trifluoroacetic acid at room temperature followed by neutralization with aqueous sodium hydroxide affords the desired anilines 5. The requisite carbamic acid esters 6, wherein R_9 - R_{12} and G are described as above, are prepared as shown in Method 2C by reaction of substituted acid chlorides, 8, where G is described as above, and 4-aminophenylcarbamic acid tert-butyl esters 7, wherein

 R_9 - R_{12} are described above, in the presence of triethylamine in an appropriate solvent such as dichloromethane, dimethylsulfoxide, or dimethylformamide or mixturestthereof. Carboxylic acid chlorides 8 are either commercially available or prepared from the corresponding carboxylic acid by reaction with oxalyl chloride in a suitable solvent such as dichloromethane at room temperature.

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Method 2C, 3A

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Alternatively, carbamic acid esters 6, wherein R₉-R₁₂ and G are described as above, are prepared as shown in Method 2E by reaction of substituted carboxylic acids 8a, wherein G is described as above, and an appropriately substituted 4-aminophenyl carbamic acid tert-butyl esters 7 in the presence of a suitable coupling agent such as benzotriazole-1-yloxy-tris-(dimethylamino)phosphonium hexafluorophosphate, 2-(1H-benzotriazole-1-yloxy)-1,1,3,3-tetramethyluronium hexafluorophosphage, dicyclo-hexyl carbodiimide or the like and in the presence of a tertiary amine base such as triethylamine or diisopropylethylamine in a suitable solvent such as dichloromethane, dimethylformamide and the like, at room temperature to produce the corresponding arylaminoamide.

Carboxylic acids 8a are either commercially available or are prepared according to literature methods. For example, when G is a substituted thiadiazole, the acid is available from the corresponding carboxylic acid ester by reaction with an appropriate base such as sodium or potassium hydroxide in a suitable solvent mixture such as methanol or ethanol and water at room temperature.

Similarly, when G is either substituted or unsubstituted thiazole, substituted or unsubstituted oxazole, substituted or unsubstituted isoxazole, substituted or unsubstituted isoxazole, when not commercially available, the corresponding carboxylic acid 8a is available from the corresponding ethyl or methyl ester by reaction with an appropriate base such as sodium or potassium hydroxide in a suitable solvent mixture such as methanol or ethanol and water at room temperature. These esters are either commercially available or can be prepared according to literature methods.

When the carboxylic acid ester precursors which are ultimately converted to acids 8a are not commmercially available, they may be prepared by methods known in the literature. For example, 5-substituted-1,2,3-thiadiazole-4 carboxylic acid esters may be prepared essentially according to the procedure of Caron, M J. Org. Chem. 51, 4075 (1986) and Taber, D. F., Ruckle, R. E. J. Amer. Chem. Soc. 108, 7686 (1986). Thus, according to Method 21, treatment of a beta-keto carboxylic acid ester with 4-methylbenzenesulfonyl azide or methanesulfonyl azide or the like in the presence of a tertiary amine base such triethylamine or diisopropylethylamine in a suitable solvent such as acetonitrile affords the corresponding diazo-beta-keto carboxylic acid ester. Treatment of this compound with 2,4-bis(4-methoxyphenyl)-1,3-dithia-2,4-diphosphetane-2,4-disulfide in a suitable solvent such as benzene or toluene or the like at temperatures ranging from room temperature to the reflux temperature of the solvent gives the desired 5-substituted-1,2,3-thiadiazole-4-carboxylic acid ester.

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Alternatively, 4-substituted-1,2,3-thiadiazole -5-carboxylic acid esters may be prepared essentially according to the procedure of Shafiee, A., Lalezari, I., Yazdani, S., Shahbazian, F. M., Partovi, T. J. Pharmaceutical Sci. 65, 304 (1976). Thus, according to Method 22 and 23, reaction of an appropriately substituted beta-keto carboxylic acid ester in a suitable alcoholic solvent such as methanol or ethanol with an aqueous solution semicarbazide hydrochloride at temperatures ranging from room temperature to the reflux temperature of the solvent in the presence of a suitable base such as pyridine gives corresponding semicarbazone derivative. Treatment of this compound with neat thionyl chloride at 0°C followed by treatment with an excess aqueous solution of sodium bicarbonate affords the corresponding 4-substituted-1,2,3-thiadiazole -5-carboxylic acid esters.

4-carboalkoxythiazoles are prepared essentially according to the procedure of Schöllkopf, U., Porsch, P., Lau, H. *Liebigs Ann. Chem.* 1444 (1979). Thus, according to Method 55 and 56, reaction of ethyl isocyanoacetate with N,N-dimethylformamide dimethyl acetal in a suitable alcoholic solvent such as ethanol at room temperature gives the corresponding 3-dimethylamino-2-isocyano-acrylic acid ethyl ester. A solution of this compound in a suitable solvent such as tetrahydrofuran is treated with gaseous hydrogen sulfide in the presence of a suitable tertiary amine

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base such as triethylamine or diiso-propylethylamine or the like at room temperature to give the corresponding 4-carboethoxy-thiazole.

Additional appropriately substituted thiazoles may be prepared essentially according to the procedure of Bredenkamp, M. W., Holzafel, C. W., van Zyl, W. J. Synthetic Comm. 20, 2235 (1990). Appropriate unsaturated oxazoles are prepared essentially according to the procedure of Henneke, K. H., Schöllkopf, U., Neudecker, T. Liebigs Ann. Chem. 1979 (1979). Substituted oxazoles may be prepared essentially according to the procedures of Galeotti, N., Montagne, C., Poncet, J., Jouin, P. Tetrahedron Lett. 33, 2807, (1992) and Shin, C., Okumura, K., Ito, A., Nakamura, Y. Chemistry Lett. 1305, (1994).

The following specific examples are illustrative, but are not meant to be limiting of the present invention.

EXAMPLE 1 (METHOD 1A)

4-Methoxy-3-trifluoromethyl- phenylamine

A suspension of 4-methoxy-3-trifluoromethyl-nitrobenzene (2.2 g) and iron powder (1.68 g) in ethanol (35 mL) and water (15 mL) is treated with a solution of concentrated hydrochloric acid (0.42 mL) in ethanol (6 mL) and water (3 mL) and the mixture is heated to reflux for approximately 1 hour. The mixture is then cooled, filtered, and concentrated under reduced pressure. The resulting oil is dissolved in ethyl acetate and extracted three times with 5% aqueous hydrochloric acid. The pooled acidic extracts are then cooled in an ice bath and basified with solid potassium carbonate, then extracted with ethyl acetate. These organic extracts are washed with saturated aqueous sodium chloride, dried over anhydrous sodium sulfate, concentrated under reduced pressure, then passed through a short column of silica gel (ethyl acetate is used as the eluant) to provide the desired compound as an amber oil.

Using the above procedure and appropriate starting materials the following compounds were prepared:

2,6-Dichloro-benzene-1,4-diamine

- 3-Chloro-4-methylsulfanyl-phenylamine
- 2,6-Dibromo-benzene-1,4-diamine
- 3-Chloro-4-trifluoromethyl-phenylamine
- 3-Chloro-4-ethylsulfanyl-phenylamine
- 4-Methoxy-3-trifluoromethyl-phenylamine
- 3,5-Dichloro-4-methoxy-2-methyl-phenylamine
- 5-Chloro-2-ethoxy-4-methoxy-phenylamine
- 5-Chloro-4-ethoxy-2-methoxy-phenylamine
- 5-Iodo-2,4-dimethoxy-phenylamine
- 3,5-Diiodo-2,4-dimethoxy-phenylamine
- 3,5-Dibromo-2,4-dimethoxy-phenylamine
- 5-Chloro-2-methoxy-4-methyl-phenylamine
- 2-Chloro-N(1),N(1)-dimethyl-benzene-1,4-diamine
- 3-Chloro-4-piperidin-1-yl-phenylamine
- 3-Chloro-4-pyrrolidin-1-yl-phenylamine
- N(1)-Benzyl-2-chloro-benzene-1,4-diamine
- 3-Chloro-4-(4-methyl-piperazin-1-yl)-phenylamine
- 2-Chloro-N(1)-methyl-N(1)-(1-methyl-piperidin-4-yl)-benzene-1,4-diamine
- 2-Chloro-N(1)-methyl-N(1)-(1-methyl-pyrrolidin-3-yl)-benzene-1,4-diamine
- 2-Chloro-N(1)-methyl-N(1)-phenyl-benzene-1,4-diamine
- N(1)-(1-Benzyl-pyrrolidin-3-yl)-2-chloro-N(1)-methyl-benzene-1,4-diamine
- 2-Chloro-N(1)-cyclopentyl-N(1)-methyl-benzene-1,4-diamine
- 2-[(4-Amino-2-chloro-phenyl)-(2-hydroxy-ethyl)-amino]-ethanol
- 2-Chloro-N(1)-hexyl-N(1)-methyl-benzene-1,4-diamine
- 2-Chloro-N(1)-isobutyl-N(1)-methyl-benzene-1,4-diamine
- 2-[(4-Amino-2-chloro-phenyl)-methyl-amino]-ethanol
- 2-Chloro-N(1)-(3-dimethylamino-propyl)-N(1)-methyl-benzene-1,4-diamine
- $\hbox{2-Chloro-N(1)-(2-dimethylamino-ethyl)-N(1)-methyl-benzene-1,4-diamine}\\$
- 2-Chloro-N(1)-(2-dimethylamino-ethyl)-benzene-1,4-diamine
- N(1)-(1-Benzyl-piperidin-4-yl)-2-chloro-benzene-1,4-diamine
- 2-Chloro-N(1)-(2-methoxy-ethyl)-N(1)-methyl-benzene-1,4-diamine
- 2-Chloro-N(1)-(3-dimethylamino-propyl)-benzene-1,4-diamine

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- N(1)-(1-Benzyl-pyrrolidin-3-yl)-2-chloro-benzene-1,4-diamine
- 3-Chloro-4-(1-methyl-piperidin-4-yloxy)-phenylamine
- 3-Chloro-4-(2-dimethylamino-ethoxy)-phenylamine
- 3-Chloro-4-(3-dimethylamino-propoxy)-phenylamine
- 3-Chloro-4-(1-methyl-pyrrolidin-3-yloxy)-phenylamine
- 3-Chloro-4-cyclohexyloxy-phenylamine

EXAMPLE 2 (METHOD 1B)

4-Bromo-2,4-dimethoxy-phenylamine

- A suspension of 4-bromo-2,4-dimethoxy-nitrobenzene (0.48 g) and iron powder (0.42 g) in acetic acid (10 mL) and ethanol (10 mL) is heated to 120 °C for approximately 5 hours. The mixture is then cooled, filtered, and concentrated under reduced pressure. Water is added and the mixture is cooled in an ice bath and neutralized with solid potassium carbonate and then extracted with dichloromethane.
- These organic extracts are washed with saturated aqueous sodium chloride, dried over 10 anhydrous sodium sulfate, concentrated under reduced pressure, then chromatographed over silica gel (20% ethyl acetate in hexanes is used as the eluant) to provide the desired compound as an amber oil.

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EXAMPLE 3 (METHOD 1C)

(4-Amino-2,6-dichloro-phenoxy)-acetic acid tert-butyl ester

A soution of (4-nitro-2,6-dichloro-phenoxy)-acetic acid tert-butyl ester (1 g) in ethanol (17 mL) and water (8.6 mL) is treated with iron powder (0.861 g) and ammonium chloride (86 mg) and the mixture is heated to reflux for approximately 1 20 hour. The mixture is then filtered and concentrated under reduced pressure. The resulting oil is partitioned between water and ethyl acetate, and the organic phase is then washed with saturated aqueous sodium chloride, dried over anhydrous sodium sulfate, and concentrated under reduced pressure to provide the desired compound as a pale yellow solid.

Using the above procedure and appropriate starting materials the following compounds were prepared:

4-Chloro-benzene-1,2-diamine

N-(4-Amino-2-chloro-phenyl)-acetamide

(4-Amino-2,6-dichloro-phenoxy)-acetonitrile

(4-Amino-2,6-dichloro-phenoxy)-acetic acid tert-butyl ester

(2-Amino-4-chloro-5-methoxy-phenoxy)-acetonitrile

(4-Amino-2-chloro-5-methoxy-phenoxy)-acetic acid methyl ester

(4-Amino-2-chloro-5-methoxy-phenoxy)-acetic acid tert-butyl ester

(2-Amino-4-chloro-5-methoxy-phenoxy)-acetic acid tert-butyl ester

N(1)-Benzyl-4-chloro-5-methoxy-benzene-1,2-diamine

N-(4-Amino-2-chloro-phenyl)-2-fluoro-benzamide

N-(4-Amino-5-chloro-2-hydroxy-phenyl)-acetamide

N-(4-Amino-5-chloro-2-hydroxy-phenyl)-2-fluoro-benzamide

Furan-2-carboxylic acid (4-amino-2-chloro-phenyl)-amide

(4-Amino-2-chloro-phenyl)-carbamic acid ethyl ester

N-(4-Amino-5-chloro-2-methyl-phenyl)-acetamide

N-(4-Amino-5-chloro-2-methyl-phenyl)-2-fluoro-benzamide

Furan-2-carboxylic acid (4-amino-5-chloro-2-methyl-phenyl)amide

N-(4-Amino-3-chloro-phenyl)-2-fluoro-benzamide

Furan-2-carboxylic acid (4-amino-3-chloro-phenyl)-amide

N-(4-Amino-2-chloro-phenyl)-2-dimethylamino-acetamide

N-(4-Amino-2-chloro-phenyl)-2-piperidin-1-yl-acetamide

N-(4-Amino-2-chloro-phenyl)-2-morpholin-4-yl-acetamide

N-(4-Amino-2-chloro-phenyl)-methanesulfonamide

N-(4-Amino-2-chloro-phenyl)-benzamide

N-(4-Amino-2-chloro-phenyl)-2-diethylamino-acetamide

N-(4-Amino-2-chloro-phenyl)-2-pyrrolidin-1-yl-acetamide

N-(4-Amino-2-chloro-phenyl)-2-azepan-1-yl-acetamide

N-(4-Amino-2-chloro-phenyl)-2-(2-methyl-piperidin-1-yl)-acetamide

N-(4-Amino-2-chloro-phenyl)-2-(3-methyl-piperidin-1-yl)-acetamide

- 3-Chloro-benzene-1,2-diamine
- 4-Chloro-N,N-dimethyl-benzene-1,2-diamine

EXAMPLE 4 (METHOD 1D) 3,5-Dichloro-4-phenoxy-phenylamine

To a slurry of 3,5-dichloro-4-phenoxy-nitrobenzene (6.1 g) and tin powder (12 g) is added dropwise concentrated hydrochloric acid (60 mL). Ethanol (60mL) is added and the mixture is heated to reflux for approximately 1 hour. The mixture is then cooled in an ice bath and basified by addition of solid sodium hydroxide. The resulting suspension is filtered through a pad of diatomaceous earth and extracted three times with ethyl acetate. The combined organic extracts are then washed with saturated aqueous sodium chloride, dried over anhydrous magnesium sulfate, and concentrated under reduced pressure to provide the desired product as a yellow solid. Recrystallization from ethyl acetate-hexanes provided the product as a pale yellow solid.

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Using the above procedure and appropriate starting materials the following compounds were prepared:

- 1-Furan-2-yl-ethylamine
- 3-Chloro-4-isopropoxy-phenylamine
- 2-Butoxy-5-chloro-4-methoxy-phenylamine
- 3,5-Dichloro-2-methoxy-4-methyl-phenylamine
- 2-Benzyloxy-5-chloro-4-methoxy-phenylamine
- 4-Benzyloxy-5-chloro-2-methoxy-phenylamine
- 5-Fluoro-2,4-dimethoxy-phenylamine
- (4-Amino-2,6-dichloro-phenoxy)-acetic acid ethyl ester
- 3,5-Dichloro-4-phenoxy-phenylamine
- 2-(4-Amino-2-chloro-5-methoxy-phenoxy)-acetamide
- (4-Amino-2-chloro-5-methoxy-phenoxy)-acetonitrile
- 2-(2-Amino-4-chloro-5-methoxy-phenoxy)-ethanol

- 2-(4-Amino-2-chloro-5-methoxy-phenoxy)-ethanol
- 4-(4-Amino-2-chloro-5-methoxy-phenoxy)-butyronitrile
- 4-Amino-2-chloro-5-methoxy-phenol
- 2-Amino-4-chloro-5-methoxy-phenol
- 5-Chloro-4-methoxy-2-morpholin-4-yl-phenylamine
- 4-Chloro-5-methoxy-N(1),N(1)-dimethyl-benzene-1,2-diamine
- 5-Chloro-4-methoxy-2-piperidin-1-yl-phenylamine
- 5-Chloro-4-methoxy-2-pyrrolidin-1-yl-phenylamine
- 2-Chloro-N(1)-cyclohexyl-N(1)-methyl-benzene-1,4-diamine
- N(2)-Benzyl-4-methoxy-benzene-1,2-diamine
- 2-(4-Amino-2-chloro-phenoxy)-ethanol
- 2-Chloro-N(1)-cyclohexyl-N(1)-ethyl-benzene-1,4-diamine
- 4-Butoxy-3-chloro-phenylamine
- (4-Amino-2-chloro-phenoxy)-acetonitrile
- 2-Chloro-N(1)-cyclohexyl-benzene-1,4-diamine
- 2-Chloro-N(1),N(1)-dipropyl-benzene-1,4-diamine
- 3-Chloro-4-(2,2,2-trifluoro-ethoxy)-phenylamine
- 3-Chloro-4-(octahydro-quinolin-1-yl)-phenylamine
- N(1)-Allyl-2-chloro-N(1)-cyclohexyl-benzene-1,4-diamine
- N-(4-Amino-2-methoxy-5-methyl-phenyl)-2-fluoro-benzamide
- Furan-2-carboxylic acid (4-amino-2-methoxy-5-methyl-phenyl)amide
- N-(4-Amino-naphthalen-1-yl)-2-fluoro-benzamide
- 3-Chloro-N,N-dimethyl-benzene-1,2-diamine
- 3-Chloro-4-propoxy-phenylamine
- 3-Iodo-4-methoxy-phenylamine
- 3-Chloro-2,4-dimethoxy-aniline
- 3-Bromo-4-methoxy-phenylamine
- 3-Chloro-4-ethoxy-phenylamine

EXAMPLE 5 (Method 1E)

(4-Amino-phenyl)-carbamic acid isobutyl ester

To a solution of N-(4-Nitro-phenyl)-isobutyrlamide (2.0 g) in 100 mL ethylene glycol monomethyl ether (100 mL) is added 10% palladium on carbon (275 mg). The mixture is hydrogenated for 2 hours at room temperature under 30 psi of hydrogen on a Parr hydrogenation apparatus. The catalyst is then removed by filtration through diatomaceous earth and the filtrate is evaporated to dryness under reduced pressure by azeotroping three times with heptane. Trituration of the residue with heptane provides the desired product as a white solid.

Using the above procedure and appropriate starting materials the following compounds were prepared:

2-Methyl-3H-benzoimidazol-5-ylamine

N-(4-Amino-phenyl)-formamide

1H-Benzoimidazol-5-ylamine

(4-Amino-phenyl)-carbamic acid isobutyl ester

N-(4-Amino-phenyl)-isobutyramide

N-(5-Amino-pyridin-2-yl)-2-methyl-benzamide

Furan-2-carboxylic acid (5-amino-pyridin-2-yl)-amide

N-(5-Amino-pyridin-2-yl)-2-fluoro-benzamide

[6-(2,2,2-Trifluoro-acetylamino)-pyridin-3-yl]-carbamic acid tert-butyl ester

N-(5-Amino-pyridin-2-yl)-2,2,2-trifluoro-acetamide

(4-Amino-benzyl)-carbamic acid tert-butyl ester

2-(3,5-Bis-trifluoromethyl-phenyl)-ethylamine

1-tert-Butyl-1H-imidazol-2-ylamine

3-(3-Dimethylamino-propyl)-5-trifluoromethyl-phenylamine

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EXAMPLE 6 (METHOD 1F)

N-(4-Amino-2-methylphenyl)-2-fluorobenzamide

A mixture of 2-fluoro-N-(2-methyl-4-nitrophenyl)benzamide (4.55 g), cyclohexene (30 mL), ethanol (70 mL), water (30 mL) and 10% palladium on charcoal (3 g) is heated at reflux for 30 minutes. The mixture is filtered through diatomaceous earth and concentrated under reduced pressure. The resulting oil is dissolved in 50 mL of ethyl acetate and cooled at 4° C for 12 hours. Filtration provides the product as a tan solid.

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Using the above procedure and appropriate starting materials the following compounds were prepared:

N-(4-Amino-2-methyl-phenyl)-acetamide

2-Methyl-benzooxazol-6-ylamine

N-(4-Amino-3-methoxy-phenyl)-acetamide

2-Acetylamino-5-amino-benzoic acid

N-(4-Amino-phenyl)-acetamide

[4-(3-Amino-benzoylamino)-phenyl]-carbamic acid tert-butyl ester

[4-(2-Amino-benzoylamino)-phenyl]-carbamic acid tert-butyl ester

N-(4-Amino-2-cyano-phenyl)-acetamide

N-(4-Amino-2,5-dimethoxy-phenyl)-2-fluoro-benzamide

Furan-2-carboxylic acid (4-amino-2,5-dimethoxy-phenyl)-amide

N-(4-Amino-2-cyano-phenyl)-2-fluoro-benzamide

Furan-2-carboxylic acid (4-amino-2-methoxy-phenyl)-amide

N-(4-Amino-2-methoxy-phenyl)-2-fluoro-benzamide

N-(4-Amino-2-methoxy-5-methyl-phenyl)-acetamide

N-(4-Amino-2-benzoyl-phenyl)-acetamide

N-(4-Amino-2-benzoyl-phenyl)-2-fluoro-benzamide

Furan-2-carboxylic acid (4-amino-2-benzoyl-phenyl)-amide

N-(4-Amino-3-methyl-phenyl)-acetamide

N-(4-Amino-3-methyl-phenyl)-2-fluoro-benzamide

Furan-2-carboxylic acid (4-amino-3-methyl-phenyl)-amide

5-Amino-2-[(2-fluorobenzoyl)amino]-N-phenylbenzamide

Furan-2-carboxylic acid (4-amino-2-phenylcarbamoyl-phenyl)amide

N-(4-Amino-naphthalen-1-yl)-acetamide

Furan-2-carboxylic acid (4-amino-naphthalen-1-yl)-amide

N-(4-Amino-2-trifluoromethyl-phenyl)-acetamide

Furan-2-carboxylic acid (4-amino-2-cyano-phenyl)-amide

Furan-2-carboxylic acid (4-amino-2-trifluoromethyl-phenyl)-amide

N-(4-Amino-2-methyl-phenyl)-2-fluoro-benzamide

Furan-2-carboxylic acid (4-amino-2-methyl-phenyl)-amide

5-Amino-2-(2-fluoro-benzoylamino)-benzoic acid

5-Amino-2-[(furan-2-carbonyl)-amino]-benzoic acid

N-(4-Amino-2-cyano-phenyl)-2,2,2-trifluoro-acetamide

N-(4-Amino-3-methyl-phenyl)-2,6-difluoro-benzamide

N-(4-Amino-3-trifluoromethyl-phenyl)-acetamide

N-(4-Amino-3-trifluoromethyl-phenyl)-2-fluoro-benzamide

N-(4-Amino-2-trifluoromethyl-phenyl)-2,2,2-trifluoro-acetamide

N-(4-Amino-2-methoxy-phenyl)-2,2,2-trifluoro-acetamide

N-(4-Amino-2-trifluoromethyl-phenyl)-2-fluoro-N-(2-fluoro-benzoyl)-benzamide

N-(4-Amino-2-trifluoromethyl-phenyl)-2-fluoro-benzamide

EXAMPLE 7 (METHOD 1G)

N-(4-Amino-2-chlorophenyl)-2-thiomorpholino-4-yl-acetamide

A solution of N-(2-chloro-4-nitrophenyl)-2-thiomorpholino-4-yl-acetamide (3.02 g) in ethanol (200 mL) is added to a solution of sodium thiosulfate (12 g) in water (60 mL). The mixture is heated at reflux for 12 hours, cooled and poured into water. The mixture is then extracted with ethyl acetate. The ethyl acetate solution is washed twice with saturated aqueous sodium chloride, dried over anhydrous potassium carbonate, filtered through a pad of diatomaceous earth and concentrated under reduced pressure to give an oil. Toluene is added and the solution chilled to give the desired product as a light orange crystalline solid.

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Using the above procedure and appropriate starting materials the following compounds were prepared:

N-(4-Amino-2-chloro-phenyl)-2-thiomorpholin-4-yl-acetamide N-(4-Amino-2-chloro-phenyl)-2-dipropylamino-acetamide

EXAMPLE 8 (METHOD 2A)

(3-Chloro-4-iodo-phenyl)-carbamic acid tert-butyl ester

To a solution of 3-chloro-4-iodo-aniline (10 g) in tetrahydrofuran (40 mL) containing diiso-propylethylamine (6.9 mL) is added di-tert-butyl-dicarbonate (8.6 g) and the mixture is heated to reflux. After approximately 15 hours additional portions of diisopropylethylamine (6.9 mL) and di-tert-butyl-dicarbonate (21 g) is added and heating is continued for approximately 24 hours. The solution is then cooled, concentrated under reduced pressure, diluted with ethyl acetate, and washed successively three times with 5% aqueous hydrochloric acid then once with saturated aqueous sodium chloride. The solution is dried over anhydrous sodium sulfate then concentrated under reduced pressure to provide the desired crude product as a brown oil. Crystallization is induced by addition of hexanes, and the collected solid material is recrystallized from hexanes to give the desired product as a white solid.

- Using the above procedure and appropriate starting materials the following compounds were prepared:
 - N'-(4-Nitro-benzoyl)-hydrazinecarboxylic acid tert-butyl ester
 - (3-Chloro-4-iodo-phenyl)-carbamic acid tert-butyl ester
 - (4-Bromo-3-chloro-phenyl)-carbamic acid tert-butyl ester
 - (3-Chloro-4-vinyl-phenyl)-carbamic acid tert-butyl ester
 - (3-Chloro-4-methylsulfanyl-phenyl)-carbamic acid tert-butyl ester
 - (4-Amino-3-chloro-phenyl)-carbamic acid tert-butyl ester
 - (4-Chloro-2-nitro-phenyl)-carbamic acid tert-butyl ester
 - (3-tert-Butoxycarbonylamino-5-chloro-phenyl)-carbamic acid tert-butyl ester

- (4-Nitro-benzyl)-carbamic acid tert-butyl ester
- (3-Bromo-5-trifluoromethyl-phenyl)-carbamic acid tert-butyl ester
- (2-Amino-3-chloro-5-trifluoromethyl-phenyl)-carbamic acid tert-butyl ester

EXAMPLE 9 (METHOD 2B)

(3-Chloro-4-vinyl-phenyl)-carbamic acid2-trimethylsilanyl-ethyl ester

To a solution of 3-chloro-4-vinyl-phenylamine (3.4 g) in N,N-dimethylformamide (44 mL) containing diisopropylethylamine (5.8 mL) is added 1-[2-(trimethylsilyl)-ethoxycarbonyl-oxy]benzotriazole (7.1 g) and the mixture is stirred at room temperature under an atmosphere of argon for three days. The solution is then diluted with water and extracted three times with diethyl ether. The combined organic extracts are washed successively with water, saturated aqueous sodium chloride, dried over anhydrous magnesium sulfate, and concentrated under reduced pressure. The resulting residue is chromatographed over silica gel (10% ethyl acetate in hexanes is used as the eluant) to provide the desired product as a yellow oil.

EXAMPLE 10 (METHOD 2C)

[4-(2-Fluoro-benzoylamino)-phenyl]-carbamic acid tert-butyl ester

To a solution of mono-N-(t-butoxycarbonyl)-1,4-phenylenediamine (1.58 g) and triethylamine (1.50 mL) in 25 mL of dichloromethane is added o-fluorobenzoyl chloride (1.20 g). A solid formed immediately forms and is filtered and washed with fresh solvent to yield a white solid, 1.90 g.

Using the above procedure and appropriate starting materials the following compounds were prepared:

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- N-(3-Methoxy-4-nitro-phenyl)-acetamide
- N-(4-Amino-phenyl)-isobutyrlamide
- 2,2,2-Trifluoro-N-(2-methoxy-4-nitro-phenyl)-acetamide
- [4-(2-Methyl-benzoylamino)-phenyl]-carbamic acid tert-butyl ester